Subculture of fin explants: a powerful optimization of fin cell production for cryobanking purposes

Nathalie Chenais, Elisabeth Sambroni, Pierre-Yves Le Bail, Catherine Labbé

▶ **To cite this version:**

Nathalie Chenais, Elisabeth Sambroni, Pierre-Yves Le Bail, Catherine Labbé. Subculture of fin explants: a powerful optimization of fin cell production for cryobanking purposes. 10. International Congress on the Biology of Fish, Jul 2012, Madison, Wisconsin, United States. , 2012. hal-02746319

HAL Id: hal-02746319
https://hal.inrae.fr/hal-02746319

Submitted on 3 Jun 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

HYPOXIA AND BRANCHIAL BASKET DEVELOPMENT: KEEPING PACE WITH A LIMITING RESOURCE

Chapman, L.J., van der Sluijs, I., and Graham, A.

Email: Lauren.chapman@mcgill.ca

Department of Biology, McGill University, 1205 Dr. Penfield Avenue, Montreal, Quebec, H3A 1B1, Canada

Symposium: Climate Change

Presentation Type: Oral

Abstract: The mouthbrooding cichlid Pseudocrenilabrus multicolor exhibits developmental plasticity in gill size that may contribute to its persistence across a wide range of habitats. An important question is whether this plasticity is reversible during ontogeny. If the plastic response to high dissolved oxygen (DO) in early life is to produce small gills, but this is not reversible, there is a risk of encountering low DO later in life without adequate compensatory mechanisms. In a rearing experiment, P. multicolor broods were split into four groups. Group 1 was raised initially under low DO, followed by a switch to high DO after 4 months. Group 2 was raised under high DO and switched to low DO. As controls, two groups were reared under either low- or high DO. Results showed that gill development tracked the temporal shift in DO. Such developmental flexibility may be important for persistence in changing and/or novel DO environments.

IN SILICO IDENTIFICATION OF DAPHNIA PULEX MIRNAS

Chen, S., McKinney, G.J., Nichols, K.M., and Sepulveda, M.S.

Email: shuai@purdue.edu

Department of Forestry and Natural Resources, Purdue University, West Lafayette, IN 47907, USA

Symposium: Fish in a Toxic World

Presentation Type: Poster

Abstract: Daphnia pulex has long served as an important model organism. With its parthenogenetic life cycle and several well-studied environmental stressor related phynotypes, D. pulex offers various benefits for aquatic toxicological research. MicroRNAs (miRNAs) are 21-25 nucleotide small non-coding RNAs that are key regulators of gene expression. In the last few years, there is increasing evidence of miRNAs and their target genes being affected by toxicants and environmental influences, suggesting an important role of miRNAs in toxicology. However, compared with other model animals that have hundreds of miRNAs reported, there are only 45 miRNAs reported to miRBase for Daphnia pulex, not to mention the absence of several deeply conserved miRNA families (like the let-7 family). As miRNAs are highly conserved between sisters groups and with the sequence of the D. pulex genome, we are trying to use computational methods to annotate miRNAs in the D. pulex genome.

SUBCULTURE OF FIN EXPLANTS: A POWERFUL OPTIMIZATION OF FIN CELL PRODUCTION FOR CRYOBANKING PURPOSES

Chênais, N and Labbé, C

Email: Nathalie.Chenais@rennes.inra.fr

INRA UR1037 Laboratoire de Physiologie et Génomique des Poissons - Campus de Beaulieu Avenue du Général Leclerc 35000 Rennes, France

Symposium: Fish Cell Cultures

Presentation Type: Poster

Abstract: Incorporation of diploid somatic fin cells in cryobank compensates for fish egg and embryo inability to cryopreservation. The obtention of large quantity of cells derived from primary culture of fin explants is therefore of considerable importance, notably in the case of threatened species where only small fin pieces can be collected. In this study developed on goldfish, we explored whether the re-plating of fin explants would allow a second round of cell production. We demonstrated that explant re-plating indeed gave a new cell population, with better proliferation properties than cells from the first explant plating. Cells from explant re-plating had the same immunochemical epithelial pattern as cells from the first explant plating. A more specific characterization of the re-plating culture using molecular markers (transcripts level and expression pattern over culture time) will be presented. Such cell production after explant re-plating counteracts the overall poor sub-culture ability of fin cells.

THE ROLES OF IROQUOIS GENES IN ZEBRAFISH RETINAL AND CARDIAC DEVELOPMENT

Cheng, S.H., Chan W.H., Choy S.W., Lee, K.

Email: bhcheng@cityu.edu.hk

Department of Biology and Chemistry, City University of Hong Kong, 83 Tat Chee Avenue, HONG KONG, Hong Kong

Symposium: Zebra Fish

Presentation Type: Oral

Abstract: The Iroquois homeobox genes play important roles in zebrafish retinogenesis and heart development. The zebrafish Iroquois homolog irx1a is expressed during retinogenesis and knockdown of irx1a results in a retinal phenotype strikingly similar to those of sonic hedgehog (shh) mutants. Analysis of shh-GFP transgene expression in irx1a knockdown retinas revealed that irx1a is required for the propagation of shh expression through the retina. Transplantation experiments illustrated that the effects of irx1a on shh expression are both cell-autonomous and non-cell-autonomous. We also found that irx2a is expressed in the developing retina and that knockdown of irx2a results in a retinal phenotype strikingly similar to that of irx1a morphants. The expression of irx2a in retina ganglion cells was shown to be irx1a- and ath5-dependent suggesting that irx1a and ath5 are transcriptional regulators of irx2a. Furthermore, irx2a expression could rescue impaired propagation of shh waves in irx1a morphants. Together, these observations suggest that Irx2 functions downstream of irx1a to control shh expression in the retina. We proposed a novel transcriptional cascade of ath5-irx1a-irx2a in the regulation of hedgehog waves during vertebrate retinal development. We are currently exploring whether similar regulatory mechanisms are controlled by the irx genes in embryonic heart. Our preliminary data supports a necessary role of irx genes in the transcriptional control of cardiogenesis

---

10th International Congress on the Biology of Fish

Madison, Wisconsin, USA.

July 15-19, 2012

Book of Abstracts

Arranged in Alphabetical Order

by Last Name of First Author

Compiled by *Don MacKinlay*